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Date 10-31-00

PTO/SB/05 (08-00)

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UTILITY PATENT APPLICATION TRANSMITTAL

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99,069 Attorney Docket No. Raymond G. Wallace First Inventor MEDICAL IMPLANT INSERTION SYSTEM Title

(Only for new nonprovisional applications under 37 CFR 1.53(b))

Express Mail Label No. Assistant Commissioner for Patents APPLICATION ELEMENTS ADDRESS TO: Box Patent Application Washington, DC 20231 See MPEP chapter 600 concerning utility patent application contents. Fee Transmittal Form (e.g., PTO/SB/17) CD-ROM or CD-R in duplicate, large table or X (Submit an original and a duplicate for fee processing) Computer Program (Appendix) Applicant claims small entity status. 2. X 8. Nucleotide and/or Amino Acid Sequence Submission See 37 CFR 1.27. (if applicable, all necessary) 3. X Computer Readable Form (CRF) - Descriptive title of the invention b. Specification Sequence Listing on: - Cross Reference to Related Applications i. ☐ CD-ROM or CD-R (2 copies); or - Statement Regarding Fed sponsored R & D - Reference to sequence listing, a table, or a computer program listing appendix Statements verifying identity of above copies - Background of the Invention - Brief Summary of the Invention **ACCOMPANYING APPLICATION PARTS** - Brief Description of the Drawings (if filed) - Detailed Description Assignment Papers (cover sheet & document(s)) - Claim(s) Power of 37 CFR 3.73(b) Statement - Abstract of the Disclosure 10. Attorney (when there is an assignee) English Translation Document (if applicable) 4. X Drawing(s) (35 U.S.C. 113) [Total Sheets 5] Copies of IDS Information Disclosure X Citations [Total Pages [2]] 12. 5. Oath or Declaration Statement (IDS)/PTO-1449 Newly executed (original or copy) Copy from a prior application (37 CFR 1.63 (d)) **Preliminary Amendment** 13. Return Receipt Postcard (MPEP 503) (for continuation/divisional with Box 17 completed) 14. (Should be specifically itemized) DELETION OF INVENTOR(S) Certified Copy of Priority Document(s) (if foreign priority is claimed) 15. Signed statement attached deleting inventor(s) named in the prior application, see 37 CFR 16. Other: 1.63(d)(2) and 1.33(b). Application Data Sheet. See 37 CFR 1.76 17. If a CONTINUING APPLICATION, check appropriate box, and supply the requisite information below and in a preliminary amendment, or in an Application Data Sheet under 37 CFR 1.76: of prior application No.:___ Divisional Continuation-in-part (CIP) Continuation Prior application information Group / Art Unit: For CONTINUATION OR DIVISIONAL APPS only: The entire disclosure of the prior application, from which an oath or declaration is supplied under Box 5b, is considered a part of the disclosure of the accompanying continuation or divisional application and is hereby incorporated by reference. The incorporation can only be relied upon when a portion has been inadvertently omitted from the submitted application parts. 18. CORRESPONDENCE ADDRESS Customer Number or Bar Code Label Correspondence address below or X (Insert Customer No. or Attach bar code fabel here): Larry W. McKenzie Reg. No.: 28,239 Name Walker, McKenzie & Walker, P.C. #434 6363 Poplar Avenue, Address 38119 Memphis City State TNZip Code USA 901-682-648 901-685-7428 Telephone Country Fax Larry W. McKenzie 28,239 Registration No. (Attorney/Agent) Name (Print/Type) make

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TOTAL AMOUNT OF PAYMENT

(\$)	395		00
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Complete if Known				
Application Number				
Filing Date	-			
First Named Inventor	Raymond G. Wallace	i		
Examiner Name				
Group Art Unit				
Attorney Docket No.	99,069	j		

METHOD OF PAYMENT	FEE CALCULATION (continued)				
1. X The Commissioner is hereby authorized to charge indicated fees and credit any examplements to:	3. ADDITIONAL FEES				
indicated fees and credit any overpayments to: Deposit	Large Entity Small Entity Fee Fee Fee Fee Foo Bookinston				
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Deposit Walker, McKenzie &	105 130 205 65 Surcharge - late filing fee or oath				
Account Name Walker, P.C.	127 50 227 25 Surcharge - late provisional filing fee or cover sheet				
X Charge Any Additional Fee Required Under 37 CFR 1.16 and 1.17	139 130 139 130 Non-English specification				
Applicant claims small entity status.	147 2,520 147 2,520 For filing a request for ex parte reexamination				
- GBG 07 OF N 1.27	112 920* 112 920* Requesting publication of SIR prior to Examiner action				
2. X Payment Enclosed: X Check Credit card Money Order Other	113 1,840* 113 1,840* Requesting publication of SIR after Examiner action				
FEE CALCULATION	115 110 215 55 Extension for reply within first month				
1. BASIC FILING FEE	116 390 216 195 Extension for reply within second month				
Large Entity Small Entity	117 890 217 445 Extension for reply within third month				
Fee Fee Fee Fee Description	118 1,390 218 695 Extension for reply within fourth month				
Code (\$) Code (\$) Fee Paid 101 710 201 355 Utility filing fee 355	128 1,890 228 945 Extension for reply within fifth month				
, ,	119 310 219 155 Notice of Appeal				
	120 310 220 155 Filing a brief in support of an appeal				
107 490 207 245 Plant filing fee 108 710 208 355 Reissue filing fee	121 270 221 135 Request for oral hearing				
114 150 214 75 Provisional filing fee	138 1,510 138 1,510 Petition to institute a public use proceeding				
	140 110 240 55 Petition to revive - unavoidable				
SUBTOTAL (1) (\$) 355.00	141 1,240 241 620 Petition to revive - unintentional				
2. EXTRA CLAIM FEES	142 1,240 242 620 Utility issue fee (or reissue)				
Fee from Extra Claims below Fee Paid	143 440 243 220 Design issue fee				
Total Claims 11 -20** = 0 X = 0	144 600 244 300 Plant issue fee				
Independent 2 - 3** = 0 x = 0	122 130 122 130 Petitions to the Commissioner				
Multiple Dependent =	123 50 123 50 Petitions related to provisional applications				
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Large Entity Small Entity Fee Fee Fee Fee Description Code (\$) Code (\$)	581 40 581 40 Recording each patent assignment per property (times number of properties)	\$40			
103 18 203 9 Claims in excess of 20	146 710 246 355 Filing a submission after final rejection				
102 80 202 40 Independent claims in excess of 3	(37 CFR § 1.129(a)) 149 710 249 355 For each additional invention to be				
104 270 204 135 Multiple dependent claim, if not paid	149 710 249 355 For each additional invention to be examined (37 CFR § 1.129(b))				
109 80 209 40 ** Reissue independent claims over original patent	179 710 279 355 Request for Continued Examination (RCE)				
110 18 210 9 ** Reissue claims in excess of 20 and over original patent	169 900 169 900 Request for expedited examination of a design application				
SUBTOTAL (2) (\$) 0	Other fee (specify)				
**or number previously paid, if greater; For Reissues, see above	Reduced by Basic Filing Fee Paid SUBTOTAL (3) (\$) 40				
SUBMITTED BY	Complete (if applicable)				

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Larry W. McKenzie

Registration No. (Attorney/Agent)

Telephone

Date

901-685-7428

10-31-00

28,239

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SPECIFICATION

(3) TITLE OF THE INVENTION:

MEDICAL IMPLANT INSERTION SYSTEM

- (4) CROSS-REFERENCE TO RELATED APPLICATIONS:
- 5 Not Applicable.
 - (5) STATEMENT RE FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT:

 Not Applicable.
 - (6) REFERENCE TO A "MICROFICHE APPENDIX":

Not Applicable.

(7) BACKGROUND OF THE INVENTION:

- 1. Field of the Invention: The present invention relates, in general, to medical implants such as punctal occluders or the like, and, more specifically, to systems including both medical implants and medical implant insertion instruments.
- 2. Information Disclosure Statement: Various small medical implants such as
 15 myringotomy tubes, punctal occluders (punctum plugs), and the like are often sold pre-loaded on disposable insertion instruments as a sterile unit or kit. Such practices save implantation time and insure that the implants are offered for implantation in a sterile condition.

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Punctal occlusion is becoming the most accepted clinical treatment for dry eye and related conditions. Today, all known suppliers of punctal occluders (punctum plugs) sell their plugs pre-loaded on insertion instruments as a sterile unit or kit (one sterile insertion instrument per sterile punctum plug). When the insertion of the plug is complete, the entire insertion instrument is immediately discarded. Unfortunately, this results in the majority of the purchase price of the punctum plug kit being discarded. This wasteful disposal of the entire insertion instrument has resulted in an artificially high delivery cost of punctal occlusion, a very inefficient use of valuable resources and a very unfortunate contribution to non-degradable waste in our environment.

// A preliminary patentability search conducted in class 606, subclasses 108, 109, 185 and 191 produced the following patents which appear to be relevant to the present invention:

Akiyama, U.S. Patent 3,888,258, issued June 10, 1975, discloses an apparatus for introducing a drain for the eardrum.

Garnett et al., U.S. Patent 3,897,786, issued August 5, 1975, discloses a disposable apparatus for inserting a myringotomy tube.

Walchle et al., U.S. Patent 3,913,584, issued October 21, 1975, discloses an otological vent tube inserter.

Darnell, U.S. Patent 4,473,073, issued September 25, 1984, discloses a myringotomy tube inserter.

Leigh, U.S. Patent 5,172,701, issued December 22, 1992, discloses a single use biopsy device.

Arick, U.S. Patent 5,681,323, issued October 28, 1997, discloses a cricothyrotomy tube insertion device.

Mendius, U.S. Patent 5,741,292, issued April 21, 1998, discloses a punctum plug

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inserting instrument.

Wallace, U.S. Patent 5,830,171, issued November 3, 1998, discloses a punctal occluder.

Richter et al., U.S. Patent 5,868,697, issued February 9, 1999, discloses an intraocular implant and delivery device.

Nothing in the known prior art discloses or suggests the present invention. More specifically, nothing in the known prior art discloses or suggests a medical implant insertion system with a medical implant cartridge including a medical implant, a head having a first end and a second end, and a pin slidably extending through the head, the pin having a first end and a second end, the first end of the pin being located adjacent the first end of the head and being removably attached to the medical implant; the second end of the pin being positioned adjacent the second end of the head; and with a medical implant insertion instrument including a handle for removable attachment to the second end of the head of the medical implant cartridge, collet means for attachment to the second end of the pin of the medical implant cartridge when the handle is attached to the second end of the head of the medical implant cartridge, and actuator means for causing the medical implant to detach from the pin of the medical implant cartridge.

(8) BRIEF SUMMARY OF THE INVENTION:

The present invention provides a medical implant insertion system. A basic concept of the present invention is to provide a medical implant insertion system that consist, in general, of two components, a high quality reusable insertion instrument and a sterile, single use, pre-loaded cartridge.

The medical implant insertion system of the present invention comprises, in general, a medical implant cartridge including a medical implant, a head having a first end

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and a second end, and a pin slidably extending through the head, the pin having a first end located adjacent the first end of the head and removably attached to the medical implant, and having a second end positioned adjacent the second end of the head; and a medical implant insertion instrument including a handle for removable attachment to the second end of the head of the medical implant cartridge, collet means for attachment to the second end of the pin of the medical implant cartridge when the handle is attached to the second end of the head of the medical implant cartridge, and actuator means for causing the medical implant to detach from the pin of the medical implant cartridge.

One object of the present invention is to provide an economical, yet precise system for the delivery punctal occluders and the like.

(9) BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWINGS:

Fig. 1 is a side elevational view of the medical implant insertion system of the present invention.

Fig. 2 is a side elevational view similar to Fig. 1 but showing the medical implant of the medical implant insertion system of the present invention exposed and ready for implanting.

Fig. 3 is an enlarged view of a portion of Fig. 2, showing an initial step of the implantation of the medical implant.

Fig. 4 is a view similar to Fig. 3 but showing the medical implant fully implanted and being released from the medical implant insertion instrument of the medical implant insertion system of the present invention.

Fig. 5 is a view similar to Figs. 3 and 4 but showing the medical implant insertion instrument of the medical implant insertion system of the present invention fully separated from and being pulled away from the medical implant.

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Fig. 6 is a side elevational view of a medical implant cartridge of the medical implant insertion system of the present invention.

Fig. 7 is a exploded view of the medical implant cartridge, showing a removable cap thereof separated from remainder thereof.

Fig. 8 is a sectional view substantially as taken on line 8-8 of Fig. 1 on an enlarged scale and with portions thereof broken away for clarity.

Fig. 9 is a sectional view similar to Fig. 9 but showing the medical implant and the medical implant insertion instrument of the medical implant insertion system of the present invention separated from one another.

Fig. 10 is a front elevational view of the medical implant insertion instrument of the medical implant insertion system of the present invention.

Fig. 11 is an enlarged sectional view of one end of the medical implant insertion instrument shown in Fig. 10.

Fig. 12 is a sectional view similar to Fig. 11 but showing certain parts thereof in a moved position.

Fig. 13 is an exploded view of the medical implant insertion instrument of the medical implant insertion system of the present invention, with parts thereof shown in section for clarity.

Fig. 14 is a front elevational view of the medical implant insertion instrument of the medical implant insertion system of the present invention, shown with a protective cap thereon.

Fig. 15 is an exploded view of the head and pin of the medical implant cartridge of the medical implant insertion system of the present invention, with parts thereof shown in section for clarity.

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(10) DETAILED DESCRIPTION OF THE INVENTION:

A preferred embodiment of the medical implant insertion system of the present invention is shown in Figs. 1-15, and identified by the numeral 11. The medical implant insertion system 11 is designed for easy, economical and precise implantation of medical implants 13, and is especially designed for the implantation of punctal occluders (punctum plugs) such as the punctal occluder disclosed in Wallace, U.S. Patent 5,830,171, issued November 3, 1998, incorporated herein by reference. Such a medical implant 13 includes a first end 15, a second end 17, and an aperture 19 extending into the second end 17 (see Figs. 8 and 9) for receiving the tip of an insertion tool, etc.

The medical implant insertion system 11 includes at least one and preferably a plurality of medical implant cartridges 21. Each medical implant cartridge 21 incudes a medical implant 13, a head 23 having a first end 25 and a second end 27, and a pin 29 having a first end 31 and a second end 33. The pin 29 slidably extends through the head 23 with the first end 31 of the pin 29 being located adjacent the first end 25 of the head 23 and being removably attached to the medical implant 13 and with the second end 33 of the pin 29 being positioned adjacent the second end 27 of the head 23. The second end 33 of the pin 29 preferably has an enlarged portion 34 formed by a collar member or the like.

The medical implant insertion system 11 includes a medical implant insertion instrument 35. The medical implant insertion instrument 35 includes a handle 37 for removable attachment to the second end 27 of the head 23 of the medical implant cartridge 21, collet means 39 for attachment to the second end 33 of the pin 29 of the medical implant cartridge 21 when the handle 37 is attached to the second end 27 of the head 23 of the medical implant cartridge 21, and actuator means 41 for causing the medical implant 13 of the medical implant cartridge 21 to detach from the pin 29 of the

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medical implant cartridge 21.

At least the medical implant 13 and first end 25 of the head 23 of each of the medical implant cartridges 21 are preferably provided sterile, and a removable cap 43 is preferably provided for protecting the sterile medical implant 13, etc. The cap 43 is preferably a clear disposable member for being snapped over the first end 25 of the head 23 of the medical implant cartridge 21 similar to the removable cap of a fountain pen or the like.

The actuator means 41 preferably includes an actuator body 45 fixedly attached to the collet means 39 so that the collet means 39 will move with the actuator body 45, and an actuator button 47 for causing the actuator body 45 to move from a first or out position to a second or in position. The actuator means 41 also preferably includes an urging means 49, preferably a coil spring 50 or the like, for urging the actuator body 45 to the out position. The actuator body 45 preferably includes a inclined plane portion 51, and the actuator button 47 preferably includes a pusher portion 53 for engaging the inclined plane portion 51 of the actuator body 45 so that downward movement of at least one end of the actuator button 47 will cause the actuator body 45 to move to the in position. A protective pen clip style cover 54 may be provided for snapping over the first end of the handle 37 when then medical implant cartridge 21 is not mounted thereon for protecting the collet means 39, etc., and for allowing the medical implant insertion instrument 35 to be clipped to a physician's pocket similar to a fountain pen or the like.

The actual construction, design and size of the medical implant insertion system 11 may vary as will now be apparent to those skilled in the art. When used for inserting punctal occluders, the medical implant insertion system 11 is preferably substantially the

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same size and has substantially the same appearance as a typical fountain pen.

The head 23 may be constructed from two basic parts, an elongated cannula 55 and a body 57. The body 57 has a central aperture 59 therethrough sized on the first end so that the second end of the cannula 55 can be pushed thereinto to secure the cannula 55 and body 57 firmly together, and sized on the second end so that the first end of the handle 37 can be snapped thereinto to removably secure the handle 37 and medical implant cartridge 21 together. The cannula 55 and body 57 can, of course, be constructed as an integral, one-piece unit out of plastic or the like. The pin 29 may consist of an elongated metal wire 61 sized so that the first end thereof can be tightly pushed into the aperture 19 in the medical implant 13 to secure the medical implant 13 thereto, and a silicone collar 63 glued or otherwise fixed to the second end of the wire 61 to form the enlarged portion 34 of the second end 33 of the pin 29. To mount the pin 29 to the head 23, the first end 31 of the pin 29 is merely placed into the second end of the aperture 59, shook until it enters the cannula 55, and then pushed through the cannula 55 until the first end 31 of the pin 29 extends past the first end of the cannula 55. The medical implant 13 can then be placed on the first end 31 of the pin 29 and the cap 43 snapped onto the first end 25 of the body 57 of the head 23 over the medical implant 13. The entire medical implant cartridge 21 is sterilized and preferably packaged in a sterile package to allow removal of the sterile medical implant cartridge 21 using a standard "peel and drop" technique. The medical implant cartridge 21 is preferably provided as a tray having ten individually sterile, tear off packages, each including an individually sterile medical implant cartridge 21.

The handle 35 may be constructed in two anodized aluminum parts, a barrel front 65 and a barrel back 67 glued or cemented together during assembly. The barrel front 65 has a central aperture 69 that extends completely therethrough and a slot 71 that opens

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into the central aperture **69** for receiving the actuator button **47**. The first end of the aperture **69** is preferably reduced or stepped down relative to the second end of the aperture **69**. The barrel back **67** preferably has a dead end, central aperture **73** that extends rearwardly from the first end thereto.

The collet means 39 may be machined or otherwise formed with a slotted cylindrical first end having a central aperture 75 in at least the first end thereof for receiving the enlarged portion 34 of the pin 29 in a manner to hold the pin 29 to the collet means 39 for movement with the collet means 39. The central aperture 75 preferably extends completely through the collet means 39.

The actuator body 45 may be machined or otherwise formed with a boss 77 on the first end thereof for being inserted into and glued to the second end of the aperture 75 of the collet means 39 to secure the collet means 39 and actuator means 41 together. A flange 79 is preferably provided on the actuator body 45 adjacent the boss 77, and a second boss 81 is provided on the second end of the actuator body 45, with the inclined plane portion 51 located between the flange 79 and boss 81 and with the boss 81 having a cross sectional area smaller that the cross sectional area of the actuator body 45 immediately adjacent the boss 81.

To assemble the handle 37, the boss 77 of the actuator body 45 is inserted into the second end of the aperture 75 in the collet means 39 and the two parts glued together to join the collet means 39 and actuator means 41 together as a integral part. The coil spring 50, etc., is placed into the aperture 73 in the barrel back 67. The collet means 39 - actuator means 41 assembly is pushed into the aperture 69 of the barrel front 65 from the second end of the aperture 69. The flange 79 is engage the end of the stepped down portion of the aperture 69 to prevent the collet means 39 - actuator means 41 assembly from passing completely through the aperture 69. Next, the barrel

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front 65 and barrel back 67 are pushed together and glued or cemented together, etc., with the boss 81 on the second end of the actuator body 45 extending into the center of the coil spring 50, etc., to align the parts together. The slotted end 83 of the actuator button 47 is then slid into first end of the slot 71 and the rear end of the button 47 is pressed toward into the slot 71 until the button 47 snaps into place on the barrel front 65.

In the preferred manner of using the medical implant insertion system 11, a sterile package containing a sterile medical implant cartridge 21 is opened, using a standard "peel and drop" technique to drop the sterile medical implant cartridge 21 onto the physician's hand. The protective pen clip style cover 54, if used, is removed from the first end of the handle 37, and the second end 27 of the head 23 of the medical implant cartridge 21 is snapped onto the first end of the handle 37. When the second end 27 of the head 23 of the medical implant cartridge 21 is snapped onto the first end of the handle 37, the collar 63 of the pin 29 will extend into the central aperture 75 of the first end of the collet means 39. The removable cap 43 can then be gently removed from the head 23 by being pulled straight out, to expose the sterile medical implant 13 for insertion. The insertion of the medical implant 13 should follow standard or desired medical procedures. For example, in the case of a punctal occluder, dilation of the punctum and the use of topical anesthetic may or may not be required. A drop of ocular lubricant and/or antibiotic drop may be placed on the occluder to help facilitate insertion. The physician should hold the handle 37, using a natural grip, with the intended "trigger-finger" oriented over the actuator button 47. The instrument can then be used to insert the medical implant 13 to the proper position. Only after the implant 13 is in its desired position, the physician smoothly pushes the actuator button 47 to cause the actuator body 45 to move to the in position, and cause the collet means 39 to retract

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the pin 29 to the in position, separating the medical implant 13 from the pin 29, etc. Care should be taken not to prematurely push the actuator button 47 and prematurely release the implant 13. After insertion, the insertion site should be carefully inspected to confirm that the implant 13 has been properly placed. If adjustment is necessary, the use of forceps or a small dilator may be helpful. The remainder of the used medical implant cartridge 21 can then been pulled from the handle 37 and discarded, leaving the medical implant insertion instrument 35 for re-use.

Although the present invention has been described and illustrated with respect to a preferred embodiment and a preferred use therefor, it is not to be so limited since modifications and changes can be made therein which are within the full intended scope of the invention.

(11) CLAIM OR CLAIMS:

1. A medical implant insertion system comprising:
(a) a medical implant cartridge including:
a medical implant,
a head having a first end and a second end, and
a pin slidably extending through said head, said pin having a first end and
a second end, said first end of said pin being located adjacent said first end of said head
and being removably attached to said medical implant; said second end of said pin being
positioned adjacent said second end of said head; and
(b) a medical implant insertion instrument including:
a handle for removable attachment to said second end of said head of said
medical implant cartridge,
collet means for attachment to said second end of said pin of said medical
implant cartridge when said handle is attached to said second end of said head of said
medical implant cartridge, and
actuator means for causing said medical implant of the medical implant
cartridge to detach from said pin of said medical implant cartridge.
2. The medical implant insertion system of claim 1 in which said medical implant
cartridge is sterile.
3. The medical implant insertion system of claim 2 in which said medical implant of
said medical implant cartridge includes a removable cap for protecting said medical
implant.

- 4. The medical implant insertion system of claim 3 in which is included a plurality of said medical implant cartridges.
- 5. The medical implant insertion system of claim 1 in which said actuator means of said medical implant insertion instrument includes an actuator body fixedly attached to said collet means so that said collet means will move with said actuator body; and in which said actuator means of said medical implant insertion instrument includes an actuator button for causing said actuator body to move from a out position and to an in position.
 - 6. The medical implant insertion system of claim 5 in which said actuator means of said medical implant insertion instrument includes an urging means for urging said actuator body to said out position.
 - 7. The medical implant insertion system of claim 5 in which said actuator body includes a inclined plane portion; and in which said actuator button includes a pusher portion for engaging said inclined plane portion of said actuator body so that downward movement of said actuator button will cause said actuator body to move to said in position.
 - 8. The medical implant insertion system of claim 1 in which said second end of said pin of said medical implant cartridge has an enlarged portion for receipt by said collet means of said medical implant insertion instrument.

1 9. The medical implant insertion system of claim 8 in which said enlarged portion 2 of said pin of said medical implant cartridge includes a collar member. 1 10. A medical implant insertion system comprising: 2 (a) a plurality of sterile medical implant cartridges, each of said sterile medical implant cartridges including: 3 a sterile medical implant, 4 5 a head having a first end and a second end, 6 a pin slidably extending through said head, said pin having a first end and 7 a second end, said first end of said pin being located adjacent said first end of said head and being removably attached to said medical implant; said second end of said pin being 8 9 positioned adjacent said second end of said head, and 10 a removable cap for protecting said sterile medical implant; and 11 12 (b) a medical implant insertion instrument including: # 12 a handle for removable attachment to said second end of said head of one 13 14 14 15 of said medical implant cartridges, collet means for attachment to said second end of said pin of said one of said medical implant cartridges when said handle is attached to said second end of said head of said medical implant cartridge, and 16 actuator means for causing said medical implant to detach from said pin of 17 said one of said medical implant cartridges; said actuator means including an actuator 18 19 body fixedly attached to said collet means so that said collet means will move with said actuator body, an actuator button for causing said actuator body to move from a out 20 21 position and to an in position, and a spring member urging said actuator body to said out

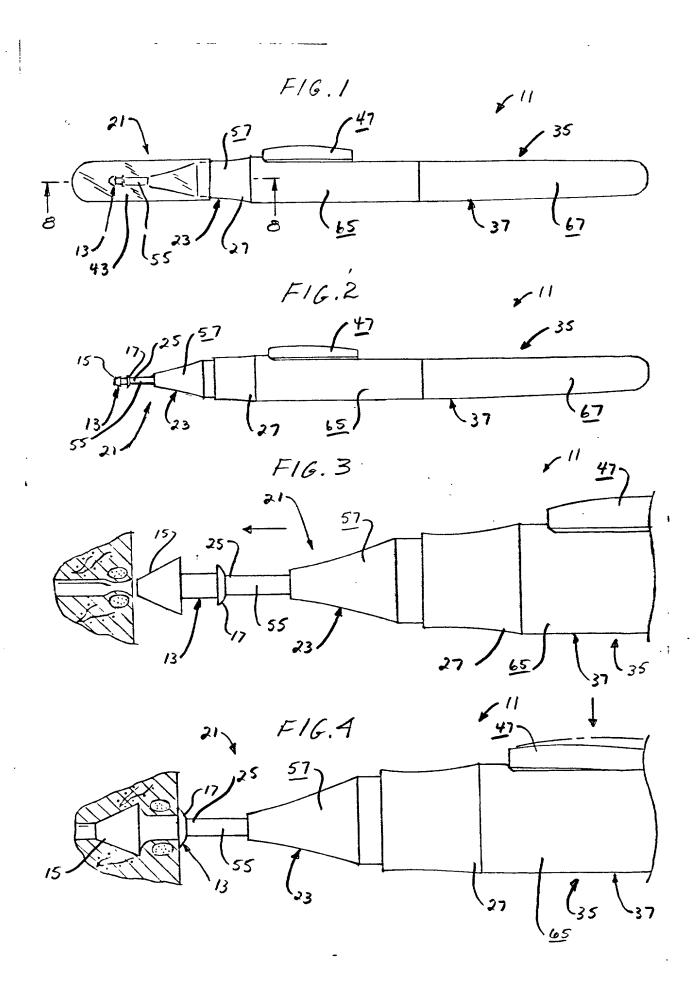
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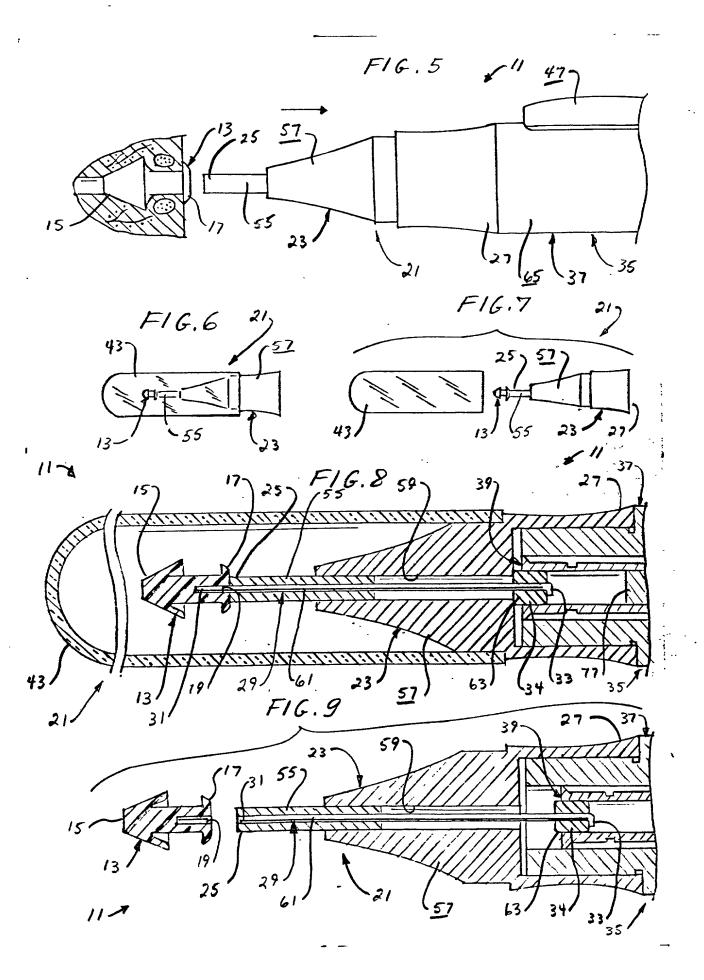
position.

- 1 11. The medical implant insertion system of claim 10 in which said actuator body
- 2 includes a inclined plane portion; and in which said actuator button includes a pusher
- 3 portion for engaging said inclined plane portion of said actuator body so that
- 4 downward movement of said actuator button will cause said actuator body to move to
- 5 said in position.

(12) ABSTRACT OF THE DISCLOSURE:

A medical implant insertion system comprising a medical implant cartridge including a medical implant, a head having a first end and a second end, and a pin slidably extending through the head, the pin having a first end and a second end, the first end of the pin being located adjacent the first end of the head and being removably attached to the medical implant; the second end of the pin being positioned adjacent the second end of the head; and a medical implant insertion instrument including a handle for removable attachment to the second end of the head of the medical implant cartridge, collet structure for attachment to the second end of the pin of the medical implant cartridge when the handle is attached to the second end of the head of the medical implant cartridge, and actuator structure for causing the medical implant to detach from the pin of the medical implant cartridge.





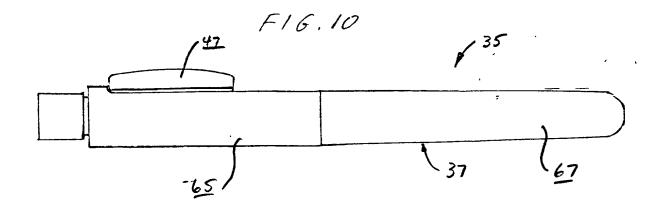
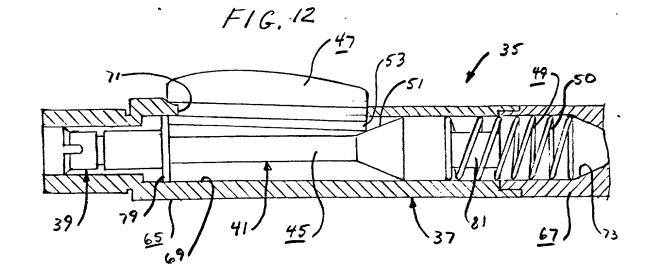
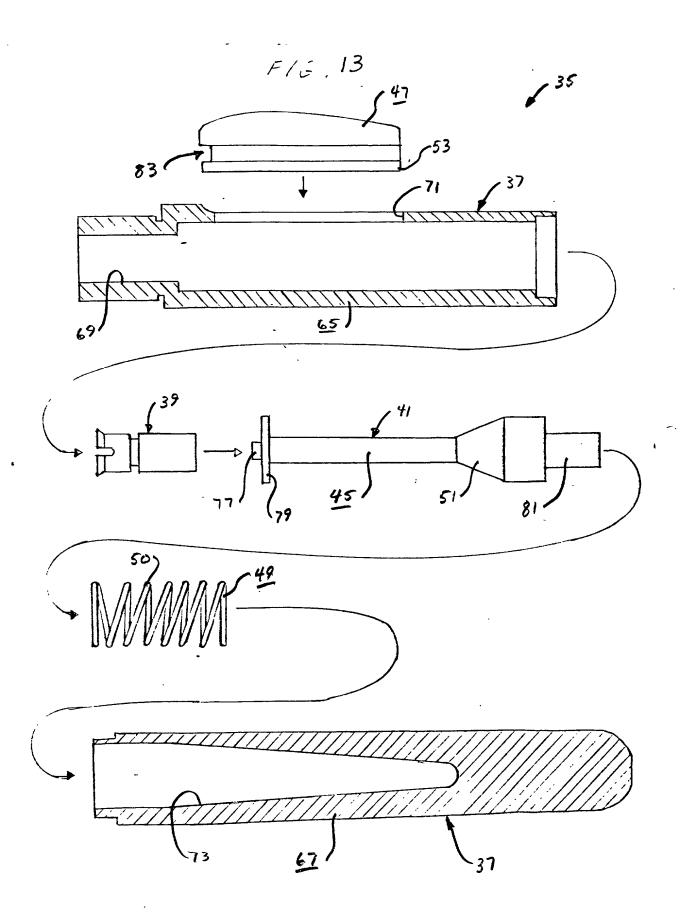


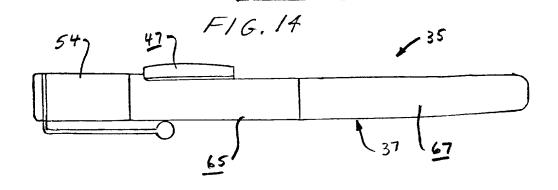
FIG.//

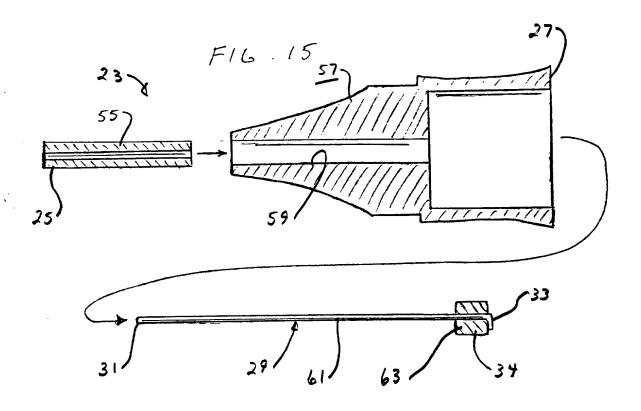
FIG.//

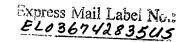
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	DECLARATION FOR UTILITY OR		Attorney Docket Number	er 99,069
DESIGN PATENT APPLICATION		First Named Inventor	Raymond G. Wallace	
		COMPLETE IF KNOWN		
	(37 CFR 1.63)		Application Number	
ſ	☐ Declaration Submitted OR With Initial Filing ☐ Declaration Submitted after Initial Filing (37 CFR 1.16 (e)) required)	Filing Date		
•		Group Art Unit		
			(37 ČFR 1.16 (e))	Examiner Name

As a below named inventor, I he	reby declare that:				
My residence, mailing address, and citizenship are as stated below next to my name.					
I believe I am the original, first and names are listed below) of the sub	I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:				
MEDICAL IMPLANT II	NSERTION SYSTE	EM			
the specification of which	(T	Title of the Invention)			
is attached hereto					
OR		as United S	States Application	Number or PCT International	
☐ was filed on (MM/DD/YYYY)				(if applicable).	
Application Number	and was a	mended on (MM/DD/YY	YY)	(ii applicable).	
I hereby state that I have reviewed amended by any amendment spec	d and understand the co	ontents of the above ide	ntified specificatio	n, including the claims, as	
I acknowledge the duty to disclose in-part applications, material inform PCT international filing date of the	continuation-in-part ap	plication.			
I hereby claim foreign priority ben- certificate, or 365(a) of any PCT in America, listed below and have certificate, or any PCT international	nternational application also identified helow	which designated at le	ast one country o	ther than the United States of	
Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached? YES NO	
			0000	0000	
Additional foreign application n					
I hereby claim the benefit under	35 U.S.C. 119(e) of an	y United States provisio	nal application(s)	listed below.	
Application Number(s)	Filing Dat	e (MM/DD/YYYY)	numbers supplem	al provisional application are listed on a ental priority data sheet 02B attached hereto.	

[Page 1 of 2]
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DECLARATION — Utility or Design Patent Application

Direct all corre		Customer Numb or Bar Code Lab				OR [X c	Correspondence address below
Name	Larry W. McKen	zie				-		
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Country	USA	Te	elephone	e 901-	-685-	-7428		Fax 901-682-6488
made are punis	I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.							
NAME OF S	OLE OR FIRST INV	ENTOR:			A petit	tion has b	een file	ed for this unsigned inventor
Given Name (first and middle	D				Family or Surr	Name	Jallac	
Inventor's Signature	Inventor's							
Residence: City	y Memphis			State TN		Country	USA	Citizenship US
Mailing Address	s 3549 Sunrise	e Lane						
Mailing Address	<u>s</u>							
City Memph	is	State TN			ZiP	38133		Country USA
NAME OF SE	ECOND INVENTOR	<u>:</u>			A petit	tion has b	een file	ed for this unsigned inventor
Given Name (first and middle	e [if any]) Gary A	٨.			Family or Surn		Tatge	
Inventor's Signature	Court a	Q. V	at	<u>.</u>				Date 10/31/00
Residence: City	y Memphis			State T	N	Country	USA	Citizenship US
Mailing Address	s 3673 Ellen D	avies Cov	√e					
Mailing Address	s							
	phis	State TN			ZIP	38133		Country USA
☐ Additional in	ventors are being named	on thesu	upplemer	ntal Addition	ial Inver	ntor(s) shee	t(s) PTO	/SB/02A attached hereto.

EL036742835US

Re: Patent Application

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of: Raymond G. Wallace and Gary A. Tatge

Assignee: Odyssey Medical, Inc..

For: MEDICAL IMPLANT INSERTION SYSTEM

Docket No.: 99,069

ASSISTANT COMMISSIONER FOR PATENTS WASHINGTON, D.C. 20231

POWER OF ATTORNEY AND 1 CERTIFICATE UNDER 37 C.F.R. § 3.73(b) 2 3 The undersigned, assignee of the entire interest in and to an application of 4 Raymond G. Wallace and Gary A. Tatge for U.S. Letters Patent for a MEDICAL 5 IMPLANT INSERTION SYSTEM, executed by the inventors on the 31st day of October, 6 2000, and further identified by Docket No. 99,069, hereby appoints the following 7 attorneys to prosecute this application and transact all business in the Patent and 8 Trademark Office in connection therewith: 9 Larry W. McKenzie Russell H. Walker Registration No. 28,239 10 Registration No. 35,401 11 Send correspondence to: 12 Walker, McKenzie & Walker, P.C. 13 6363 Poplar Ave., Suite 434 14 Memphis, Tennessee 38119-4896 15 Direct telephone calls to Larry W. McKenzie at (901) 685-7428. 16 The below-identified Assignee certifies that it is the assignee of the entire 17 right, title and interest in the provisional patent application identified above by virtue of an Assignment from the Inventor(s), a copy of which is attached hereto. 18

1	The undersigned has reviewed all the documents in the chain of title of the
2	patent application identified above and, to the best of the undersigned's
3	knowledge and belief, title is in the Assignee identified below.
4	The undersigned (whose title is supplied below) is empowered to sign this
5	certificate on behalf of the Assignee.
6	I hereby declare that all statements made herein of my own knowledge are
7	true, and that all statements made on information and belief are believed to be
8	true; and further, that these statements are made with the knowledge that willful
9	false statements, and the like so made, are punishable by fine or imprisonment, or
10	both, under Section 1001, Title 18 of the United States Code, and that such willful
11	false statements may jeopardize the validity of the application or any patent
12	issuing thereon.
13	Odyssey Medical, Inc., Assignee
14 15	Date: 10/31/00 By: Gary A. Targe President
16	riesident V